



Dublin, 01 January 2024

DECLARATION OF CONFORMITY

Regulation (EU) 2017/745 Of the European Parliament and of The Council on Medical Devices (MDR)

We,

LM Global Design Ltd.

with registered place of business in: Suite 123, The Capel Building, Mary's Abbey, Dublin 7 D07 VY68, Ireland SRN: N/A

as a legal manufacturer hereby declare under our sole responsibility, that the medical device listed below with related accessories, meets the general safety and performance requirements of Annex I and that it is in conformity with the Regulation (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL on medical devices (MDR).

The product specified below is a "technical aid for the disabled", classified as Class I, medical device. The classification is based on the requirements of annex VII & Rule I, of the MDR. Conformity assessment was carried out according to Art. 52, pt. 7 and Annex II of the MDR. The CE mark has been affixed on the product according to Annex V of the MDR.

TRADE NAME: COMMODE CHAIR LM - ITEM NUMBER: KING-CMD-00 TRADE NAME: COMMODE CHAIR W/ARMRESTS LM - ITEM NUMBER: KING-CMDA-00 TRADE NAME: COMMODE CHAIR W/BACKREST LM - ITEM NUMBER: KING-CMDB-00 TRADE NAME: COMMODE CHAIR W/BACKREST & ARMRESTS LM - ITEM NUMBER: KING-CMDBA-00

BASIC UDI-DI CODE: 539153269LMCMDFC ACCESSORY LIST: Soft Cushions (KING-CCMD2/4-00/KING-CCMDH4-00, KING-CBRX-00), Bucket and Lid (KING-BC-00)

Following harmonized norms and/or common specifications were used for conformity evaluation: EN ISO 14971:2019+A11:2021, PN-EN 1041:2010 + A1 2013, EN ISO 12182:2012, EN ISO 21856:2022, EN ISO 14001:2015, EN ISO 13485:2016+A11:2021.







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Important notice: LM Global Design Limited (LMGD) is the manager of personal data processing in accordance with the law and regulations on the protection of personal data. LMGD as the head of personal data processing determines the purpose and means of processing personal data and is responsible for keeping and using personal data in paper and/or electronic form.